BECEIVED CENTRAL FAX CENTER

DEC 1 3 2006

Appl. No. 10/805,099 (Docket 099/004)

Amdi. dated Dec. 13, 2006

Reply to Office Action of July 13, 2006

Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

Listing of Claims:

What is claimed as the invention is:

The invention claimed is:

- A method of generating a cell composition containing cardiomyocytes or cardiomyocyte
 precursor cells from primate phyripotent stem (pPS) cells obtained from a human blastocyst
 human embryonic stem (hES) cells, comprising:
 - a) initiating differentiation of the pPS hES cells in suspension culture by forming embryoid bodies;
 - b) culturing the initiated cells so that they differentiate into areas that undergo spontaneous contraction;
 - c) harvesting the differentiated cells;
 - d) separating the harvested cells into fractions according to their based on density; and
 - e) collecting combining the cell fractions containing cells that express cardiac troponin I (cTnI), cardiac troponin T (cTnT), or atrial natriuretic factor (ANF) from an endogenous gene;

thereby generating a cell composition containing cardiomyocytes or cardiomyocyte precursor cells.

- 2. The method of claim 1, wherein the embryoid bodies are plated onto a surface coated with gelatin or Matrigel®.
- The method of claim 1, wherein the cells are differentiated in the presence of a nucleotide analog that affects DNA methylation; such as 5 aza deoxy cytidine.

Appl. No. 10/805,099 (Docket 099/004) Amdt. dated Dec. 13, 2006 Reply to Office Action of July 13, 2006

- 4. The method of claim 1, wherein the cells are differentiated in a growth environment comprising a morphogen such as activin, and two or more growth factors.
- The method of claim 4, wherein the morphogen is an activin, and the growth factors include an insulin-like growth factor and a member of the TGFβ family.
- The method of claim 1, wherein the cells are differentiated in a growth environment containing about 20% serum or serum substitute.
- 7. The method of claim 1, wherein the harvested cells are separated by density centrifugation.
- 8. The method of claim 1, wherein the separating comprises distributing cells in the population according to based on their density, and collecting cells at combining cell fractions with a density between ~1.05 and ~1.075 g/mL.
- 9. The method of claim 1, further comprising culturing the collected colls combined cell fractions for at least 1 week in a medium containing a compound capable of forming a high energy phosphate bond, an acyl group carrier molecule, and a cardiomyocyte calcium channel modulator.
- 10. The method of claim 9, further comprising culturing the collected cells combined cell fractions for at least 1 week in a medium containing creatine, carnitine, or taurine.

11.-16. (Canceled)